

Refine Search

Search Results -

Terms	Documents
pyrogallol adj8 cyclic and L6	5

Database:

US Pre-Grant Publication Full-Text Database
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Search:

L7

Search History

DATE: Thursday, January 11, 2007 [Purge Queries](#) [Printable Copy](#) [Create Case](#)

<u>Set Name</u> side by side	<u>Query</u>	<u>Hit Count</u>	<u>Set Name</u> result set
<i>DB=PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=YES; OP=ADJ</i>			
<u>L7</u>	pyrogallol adj8 cyclic and l6	5	<u>L7</u>
<u>L6</u>	L5 and (treat\$7 or infect\$7)	76	<u>L6</u>
<u>L5</u>	L3 and (hiv or aids)	80	<u>L5</u>
<u>L4</u>	L3 and phenyl tetramer	1	<u>L4</u>
<u>L3</u>	L2 and cyclic	192	<u>L3</u>
<u>L2</u>	L1 and (514/\$ or 562/\$)	469	<u>L2</u>
<u>L1</u>	pyrogallol	13290	<u>L1</u>

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Search Results - Record(s) 1 through 5 of 5 returned.

☐ 1. Document ID: US 20050288228 A1

L7: Entry 1 of 5

File: PGPB

Dec 29, 2005

PGPUB-DOCUMENT-NUMBER: 20050288228

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20050288228 A1

TITLE: Bile-acid conjugates for providing sustained systemic concentrations of drugs

PUBLICATION-DATE: December 29, 2005

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Cundy, Kenneth C.	Redwood City	CA	US
Gallop, Mark A.	Los Altos	CA	US

US-CL-CURRENT: 514/12; 514/169, 514/44

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw D
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☐ 2. Document ID: US 20020142998 A1

L7: Entry 2 of 5

File: PGPB

Oct 3, 2002

PGPUB-DOCUMENT-NUMBER: 20020142998

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020142998 A1

TITLE: Bile-acid conjugates for providing sustained systemic concentrations of drugs

PUBLICATION-DATE: October 3, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Cundy, Kenneth C.	Redwood City	CA	US
Gallop, Mark A.	Los Altos	CA	US

US-CL-CURRENT: 514/169; 514/182, 552/515, 552/540

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw. De
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☐ 3. Document ID: US 20020099041 A1

L7: Entry 3 of 5

File: PGPB

Jul 25, 2002

PGPUB-DOCUMENT-NUMBER: 20020099041

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020099041 A1

TITLE: Bile-acid derived compounds for enhancing oral absorption and systemic bioavailability of drugs

PUBLICATION-DATE: July 25, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Gallop, Mark A.	Los Altos	CA	US
Cundy, Kenneth C.	Redwood City	CA	US

US-CL-CURRENT: 514/169

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw. De
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☐ 4. Document ID: US 7144877 B2

L7: Entry 4 of 5

File: USPT

Dec 5, 2006

US-PAT-NO: 7144877

DOCUMENT-IDENTIFIER: US 7144877 B2

TITLE: Bile-acid derived compounds for enhancing oral absorption and systemic bioavailability of drugs

PRIOR-PUBLICATION:

DOC-ID	DATE
US 20020099041 A1	July 25, 2002

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw. De
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☐ 5. Document ID: US 6984634 B2

L7: Entry 5 of 5

File: USPT

Jan 10, 2006

US-PAT-NO: 6984634

DOCUMENT-IDENTIFIER: US 6984634 B2

TITLE: Bile-acid conjugates for providing sustained systemic concentrations of drugs

PRIOR-PUBLICATION:

DOC-ID

US 20020142998 A1

DATE

October 3, 2002

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequence	Attachments	Claims	KM/C	Draw. De
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Terms

Documents

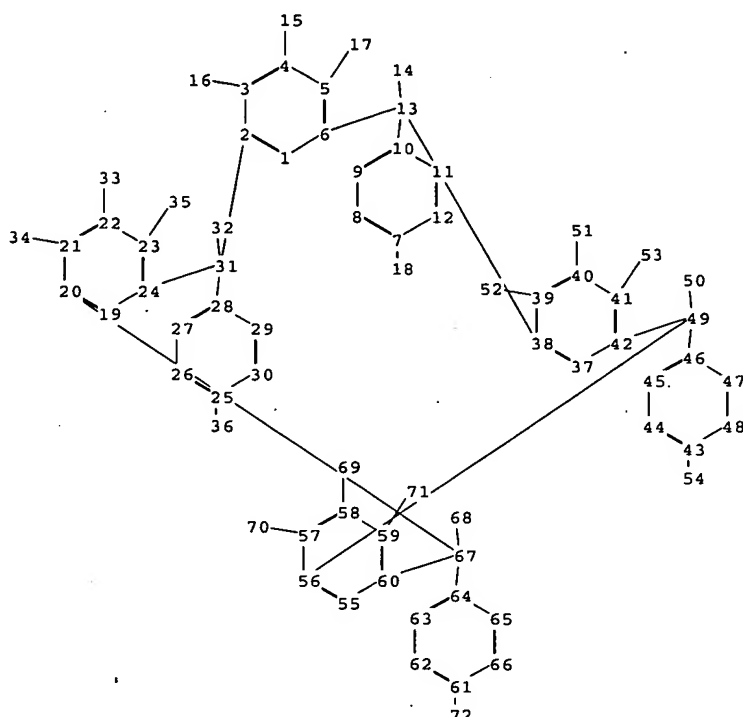
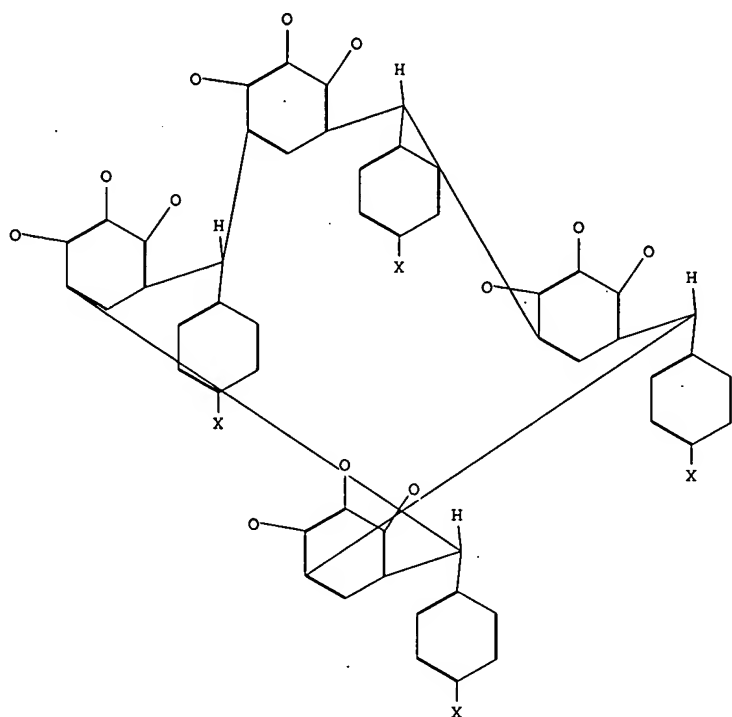
pyrogallol adj8 cyclic and L6

5

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chain nodes :

14 15 16 17 18 32 33 34 35 36 50 51 52 53 54 68 69 70 71 72

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 13 19 20 21 22 23 24 25 26 27 28 29 30 31 37 38 39
40 41 42 43 44 45 46 47 48 49 55 56 57 58 59 60 61 62 63 64 65 66 67

chain bonds :

3-16 4-15 5-17 7-18 10-13 13-14 21-34 22-33 23-35 25-36 28-31 31-32 39-52 40-51 41-53
43-54 46-49 49-50 57-70 58-69 59-71 61-72 64-67 67-68

ring bonds :

1-2 1-6 2-3 2-31 3-4 4-5 5-6 6-13 7-8 7-12 8-9 9-10 10-11 11-12 13-38 19-20 19-24 20-21
20-67 21-22 22-23 23-24 24-31 25-26 25-30 26-27 27-28 28-29 29-30 37-38 37-42 38-39 39-40
40-41 41-42 42-49 43-44 43-48 44-45 45-46 46-47 47-48 49-56 55-56 55-60 56-57 57-58 58-59
59-60 60-67 61-62 61-66 62-63 63-64 64-65 65-66

exact/norm bonds :

2-31 3-16 4-15 5-17 6-13 13-38 20-67 21-34 22-33 23-35 24-31 39-52 40-51 41-53 42-49 49-56
57-70 58-69 59-71 60-67

exact bonds :

7-18 10-13 13-14 25-36 28-31 31-32 43-54 46-49 49-50 61-72 64-67 67-68

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12 19-20 19-24 20-21 21-22 22-23
23-24 25-26 25-30 26-27 27-28 28-29 29-30 37-38 37-42 38-39 39-40 40-41 41-42 43-44 43-48
44-45 45-46 46-47 47-48 55-56 55-60 56-57 57-58 58-59 59-60 61-62 61-66 62-63 63-64 64-65
65-66

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 12:Atom
13:Atom 14:CLASS15:CLASS16:CLASS17:CLASS18:CLASS19:Atom 20:Atom 21:Atom 22:Atom 23:Atom
24:Atom 25:Atom 26:Atom 27:Atom 28:Atom 29:Atom 30:Atom 31:Atom 32:CLASS33:CLASS34:CLASS
35:CLASS36:CLASS37:Atom 38:Atom 39:Atom 40:Atom 41:Atom 42:Atom 43:Atom 44:Atom 45:Atom
46:Atom 47:Atom 48:Atom 49:Atom 50:CLASS51:CLASS52:CLASS53:CLASS54:CLASS55:Atom 56:Atom
57:Atom 58:Atom 59:Atom 60:Atom 61:Atom 62:Atom 63:Atom 64:Atom 65:Atom 66:Atom 67:Atom
68:CLASS69:CLASS70:CLASS71:CLASS72:CLASS

PASSWORD:
TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS 14 DEC 18 CA/Caplus pre-1967 chemical substance index entries enhanced
with preparation role
NEWS 15 DEC 18 CA/Caplus patent kind codes updated
NEWS 16 DEC 18 MARPAT to CA/Caplus accession number crossover limit increased
to 50,000
NEWS 17 DEC 18 MEDLINE updated in preparation for 2007 reload
NEWS 18 DEC 27 CA/Caplus enhanced with more pre-1907 records
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MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
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L1 STR

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FULL SCREEN SEARCH COMPLETED - 161 TO ITERATE

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23 ANSWERS

SEARCH TIME: 00.00.01

L2 23 SEA SSS FUL L1

L3 5 L2

=> s l3 and py<2003

22868731 PY<2003

L4 2 L3 AND PY<2003

=> d 1-2 ibib abs hitstr

L4 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2002:428850 CAPLUS

DOCUMENT NUMBER: 137:6006

TITLE: Preparation of Calixarenes as Anti-viral compounds

INVENTOR(S): Harris, Stephen J.

PATENT ASSIGNEE(S): Aids Care Pharma Limited, Ire.

SOURCE: PCT Int. Appl., 44 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002044121	A1	20020606	WO 2001-IE150	20011130 <--
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2002020992	A5	20020611	AU 2002-20992	20011130 <--
EP 1345884	A1	20030924	EP 2001-998526	20011130
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
PRIORITY APPLN. INFO.:			IE 2000-983	A 20001201
			WO 2001-IE150	W 20011130
OTHER SOURCE(S):	CASREACT 137:6006; MARPAT 137:6006			
GI				

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Title compds. I [R1 = OCH2CO2K, OCH2CO2H or OCH2CONH2; R2 = R1 or NO2; R3 = H, 2-HO2CCH2OC6H4, or 4-XC6H4 where X = halo (preferably F or Br); R4 = H or halo (preferably Br)] are prepared and disclosed as antiviral agents. Thus, II was prepared in four steps via cyclocondensation 4-fluorobenzaldehyde with pyrogallol and subsequent bromination, O-alkylation with Et bromoacetate and hydrolysis with KOH. II possessed a therapeutic index (TC50/EC50 μ m) of 4,000. I were found to have an additive effect when administered with AZT, and therefore, the compds. are useful as pharmaceutical compns. in the treatment of AIDS.

IT 433334-86-2P 433334-87-3P 433334-88-4P

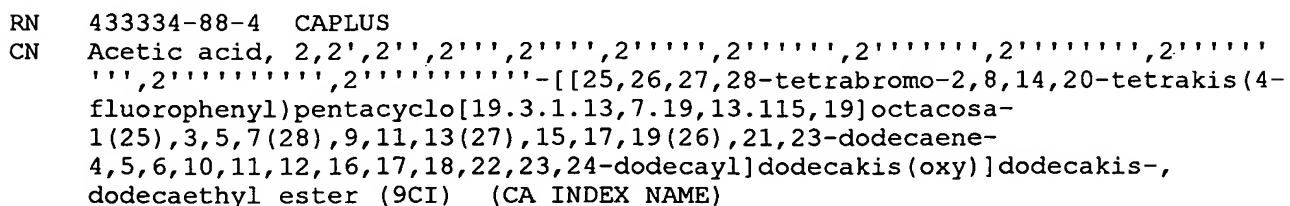
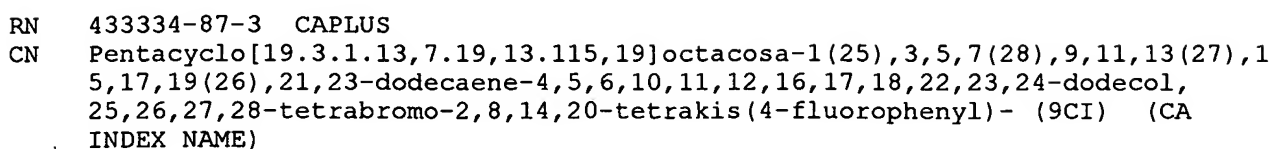
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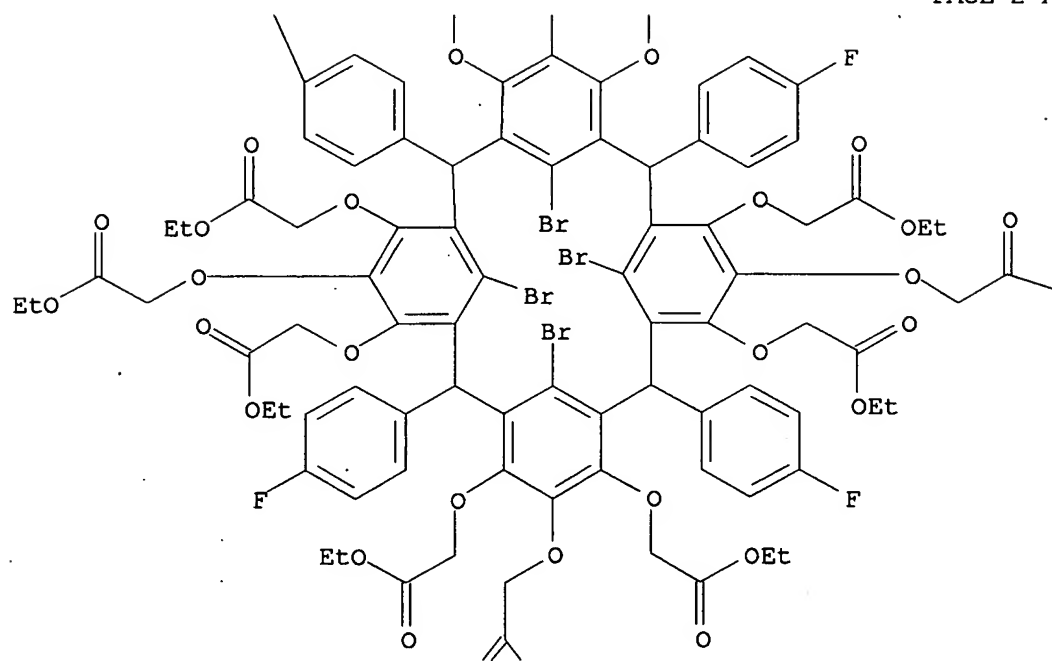
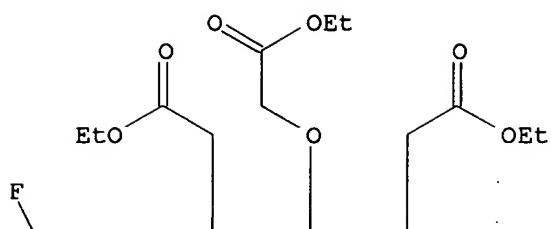
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(intermediates; preparation and antiviral activity of calixarenes as anti-AIDS agents)

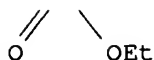
RN 433334-86-2 CAPLUS

CN Pentacyclo[19.3.1.13,7.19,13.115,19]octacos-1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecol, 2,8,14,20-tetrakis(4-fluorophenyl)- (9CI) (CA INDEX NAME)

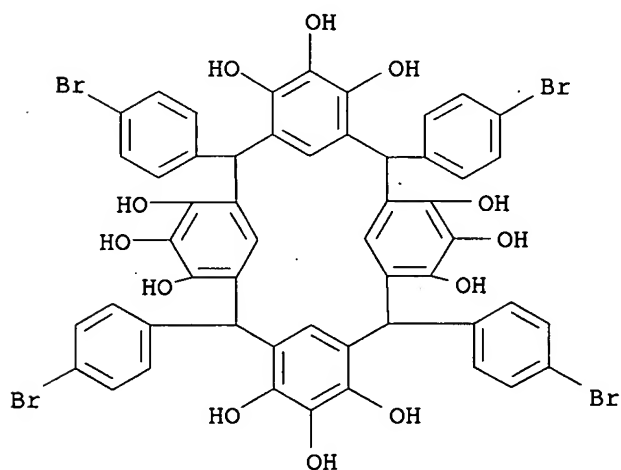




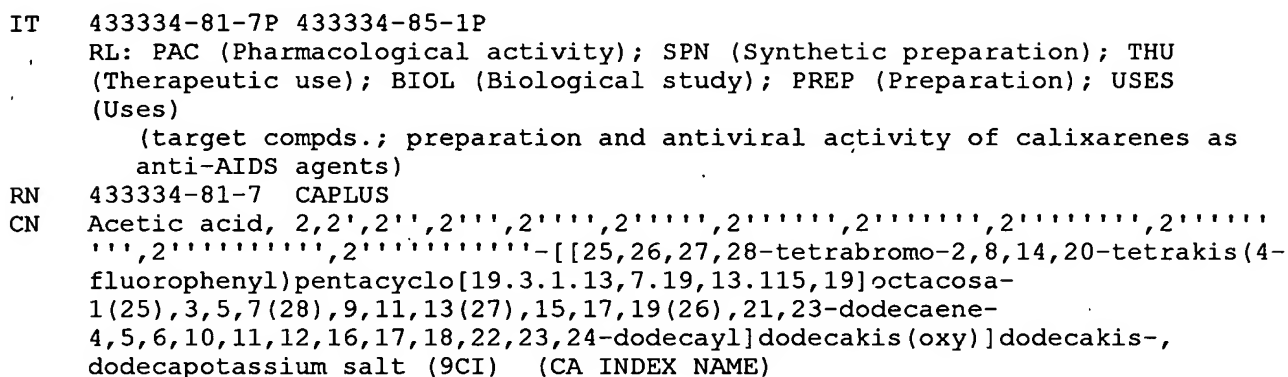
—OEt



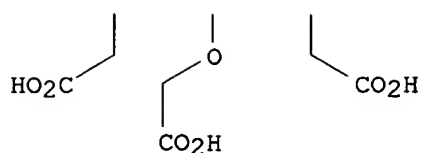
RN 433334-94-2 CAPLUS
 CN Pentacyclo[19.3.1.13,7.19,13.115,19]octacos-1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecol,2,8,14,20-tetrakis(4-bromophenyl)- (9CI) (CA INDEX NAME)



RN 433334-95-3 CAPLUS
 CN Pentacyclo[19.3.1.13,7.19,13.115,19]octacos-1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecol,25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4-bromophenyl)- (9CI) (CA INDEX NAME)



The chemical structure is a macrocycle consisting of four benzene rings connected by methylene (-CH₂-) bridges. Each of the four benzene rings has a bromine (Br) atom at the 1-position. The other three positions on each benzene ring are occupied by carboxymethoxy (-OCH₂CO₂H) groups. The macrocycle is formed by the connection of these carboxymethoxy groups, creating a large ring with four bromine atoms and eight carboxymethoxy groups.

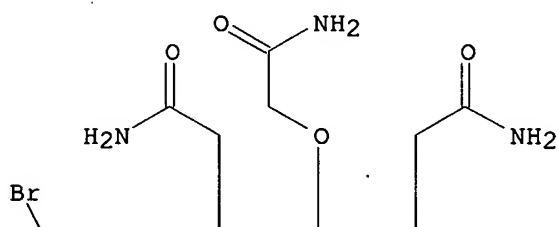


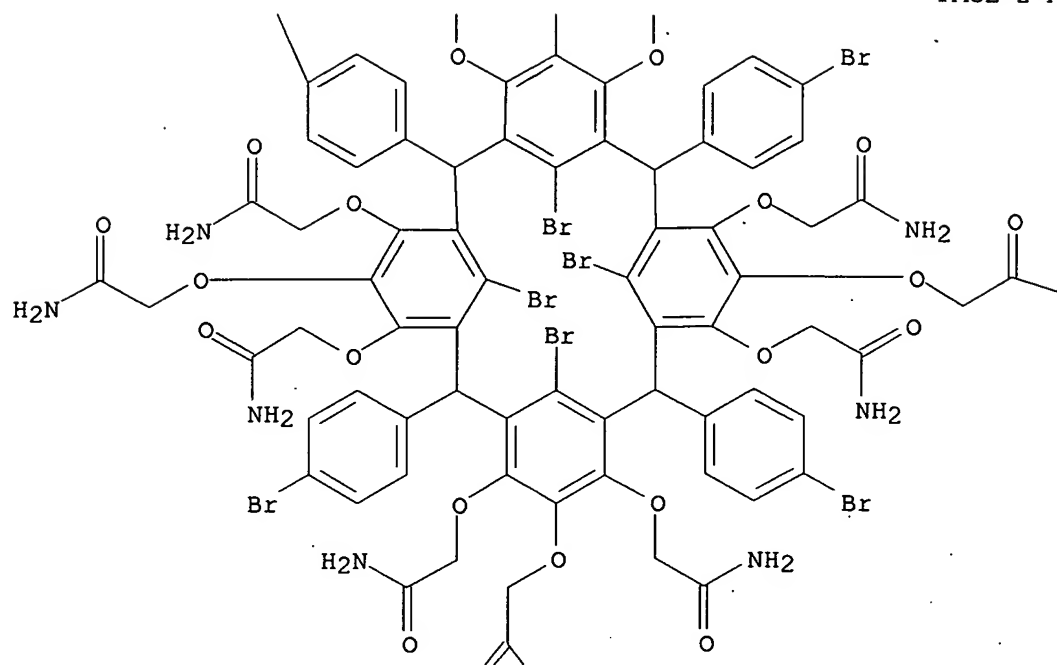
●12 K

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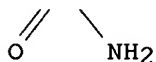
RN      433334-85-1  CAPLUS
CN      Acetamide, 2,2',2'',2''',2'''',2'''''',2''''''',2''''''',2''''''',2''''''''
        ',2''''''''''',2'''''''''''''-[25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4-
        bromophenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacosan-
        1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene-
        4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis- (9CI)
        (CA INDEX NAME)

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NH₂



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1995:994163 CAPLUS
 DOCUMENT NUMBER: 124:55584
 TITLE: Preparation of calixarene-based compounds having antibacterial, antifungal, anticancer, and anti-HIV activity
 INVENTOR(S): Harris, Stephen J.
 PATENT ASSIGNEE(S): Ire.
 SOURCE: PCT Int. Appl., 148 pp.
 CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9519974	A2	19950727	WO 1995-IE8	19950124 <--
WO 9519974	A3	19950921		
W: AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, FI, GB, HU, JP, KP, LU, NO, RO, UA, US				
RW: AT, BE, CH, DE, ES, FR, GB, GR, IE, LU, NL, SE, GA, ML, NE, SN, TD, TG				
AU 9515453	A	19950808	AU 1995-15453	19950124 <--
PRIORITY APPLN. INFO.:			IE 1994-57	A 19940124
			WO 1995-IE8	A 19950124

OTHER SOURCE(S): MARPAT 124:55584

GI For diagram(s), see printed CA Issue.

AB Calixarene-based compds., which are calixarenes or oxacalixarenes, acyclic phenyl-formaldehyde oligomers, cyclotrimeratrylene derivs., cyclic tetrameric resorcinol-aldehyde derivs. known as Hogberg compds. and cyclic tetrameric pyrogallol-aldehyde derivs., are prepared For example, calixarenes or oxacalixarenes are represented by general formula [I; $n + m = 3-8$; $m = 0-3$; $n = 0-8$; $R_1 = H$, halo, hydrocarbyl, aryl, (un)substituted hydrocarbylaryl, NO_2 , SO_3M_1 ; wherein $M_1 = \text{alkali metal}$, SO_3H ; $R_1 = OR_2$; wherein $R_2 = CH_2CO_2R_3$, CH_2CO_2Mp/p , $CH_2CONR_4R_5$; wherein $R_3 =$ (un)substituted alkyl; $M = \text{metal}$, ammonium ion; $p =$ the charge on the metal ion; R_4 or R_5 may be the same or different, or both may be part of amino acid ester of poly(amino acid ester) or one or more of the same or different amino acids or part of a cyclic polyene antibiotic/antifungal drug or part of a cyclic nitrogen heterocycle; $X = \text{halo}$, NO_2 , CO_2H , cyano, other electron withdrawing group]. Thus, n-butyraldehyde and pyrogallol in a 1:4 mixture of 37% aqueous HCl and EtOH was refluxed under N for 90 min to give a cyclic tetramer (II; $R = X = H$), which was brominated with Br in $CHCl_3$ to II ($R = H$, $X = Br$) and etherified with Et bromoacetate in the presence of K_2CO_3 in refluxing acetone to give II ($R = CH_2CO_2Et$, $X = Br$). The latter compound was saponified with KOH in refluxing EtOH, acidified with aqueous HCl, and treated with 25% aqueous NH_4OH to give II ($R = CH_2CO_2-NH_4^+$, X

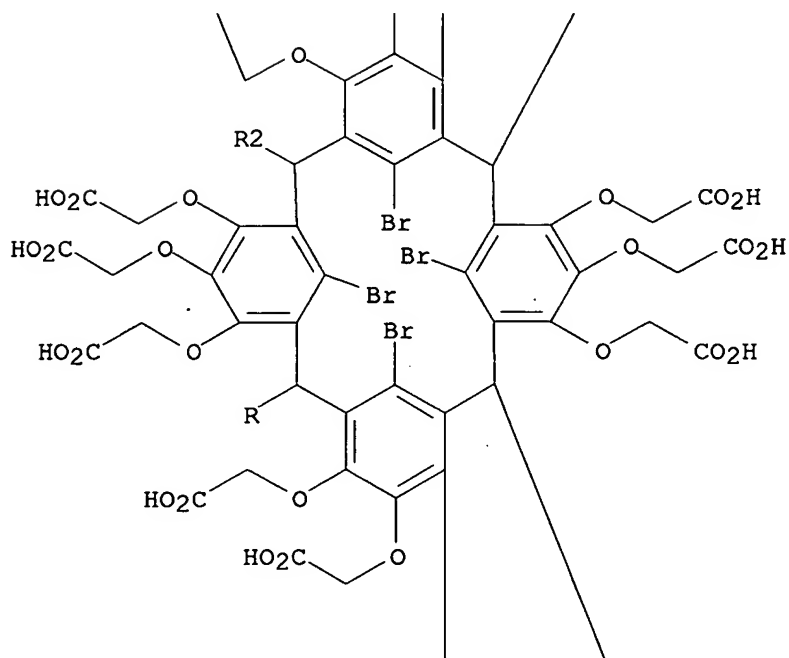
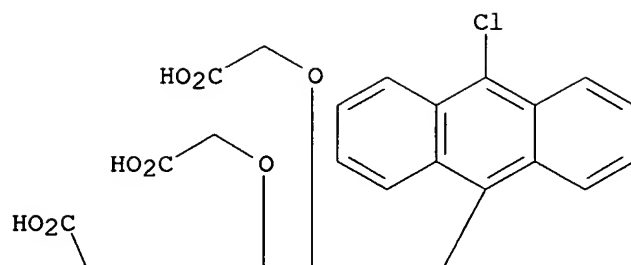
Br). The latter compound in vitro inhibited the infection of C8166 cells with HIV-2, SIV (Simian immunodeficiency virus), and HIV-1 with EC50 of 10, 20, and 0.03 μ M.

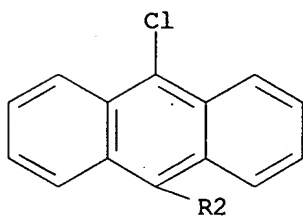
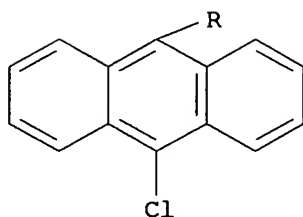
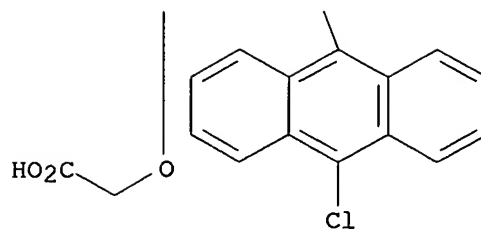
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	171799-89-6P	171799-90-9P	171799-91-0P
	171799-95-4P	171799-96-5P	171799-97-6P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of calixarene-based compds. having antibacterial, antifungal, anticancer, and anti-HIV activity)

RN 171799-80-7 CAPLUS

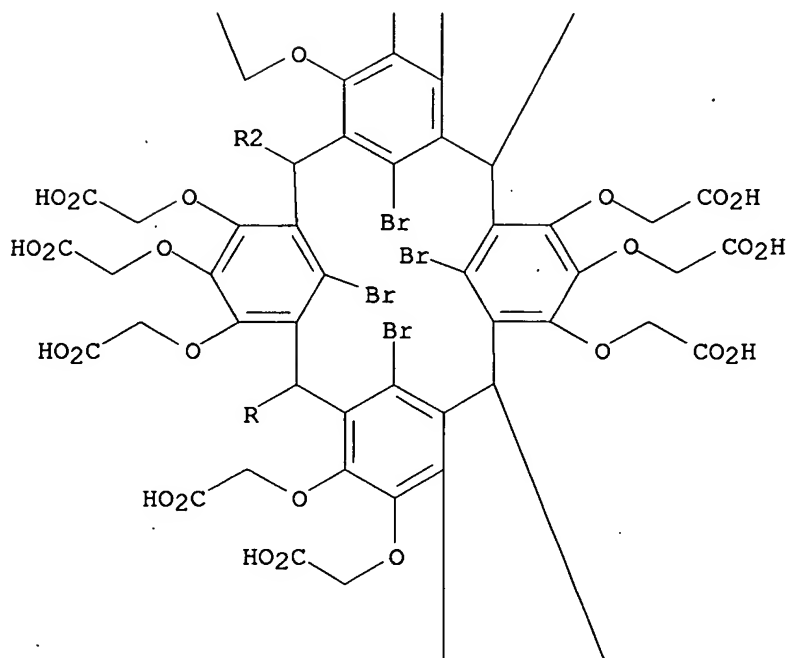
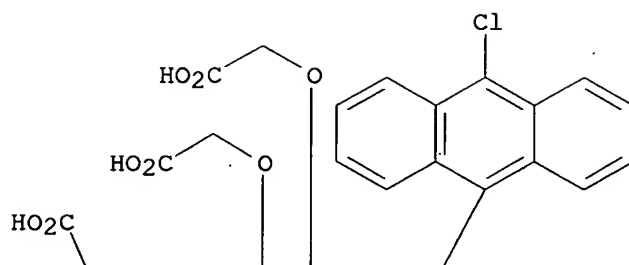
CN Acetic acid, 2,2',2'',2''',2''''',2''''',2''''',2''''',2''''',2''''',
 ''',2''''''''',2'''''''''-[25,26,27,28-tetrabromo-2,8,14,20-(10-chloro-
 9-anthracenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos-
 1(25),3,5,7(28),8,11,13(27),15,17,19(26),21,23-dodecaene-
 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis-,
 dodecapotassium salt (9CI) (CA INDEX NAME)

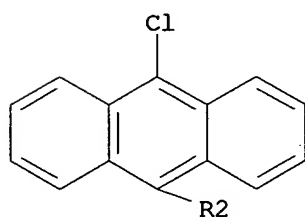
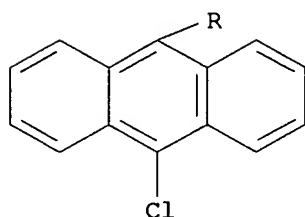
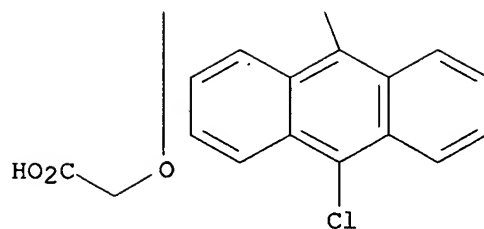




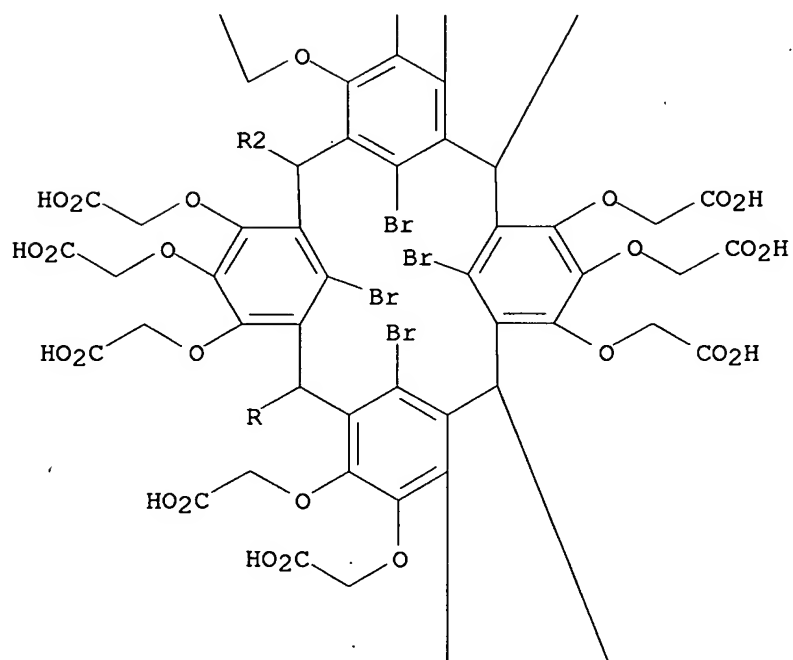
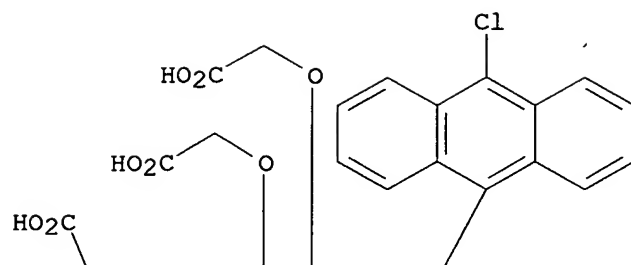
●12 K

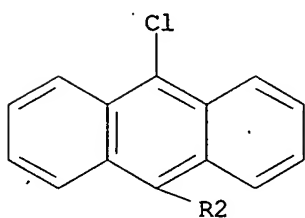
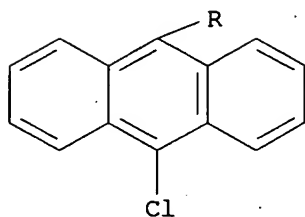
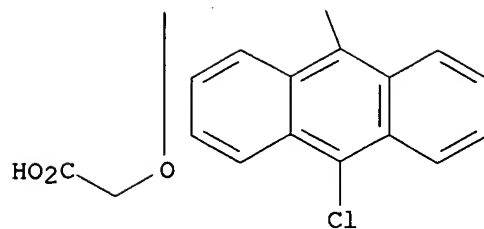
RN	171799-81-8 CAPLUS
CN	Acetic acid, 2,2',2'',2''',2'''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''''',2'''''''''''-[[25,26,27,28-tetrabromo-2,8,14,20-(10-chloro- 9-anthracenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos- 1(25),3,5,7(28),8,11,13(27),15,17,19(26),21,23-dodecaene- 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis- (9CI) (CA INDEX NAME)





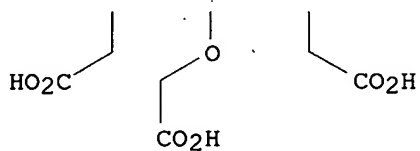
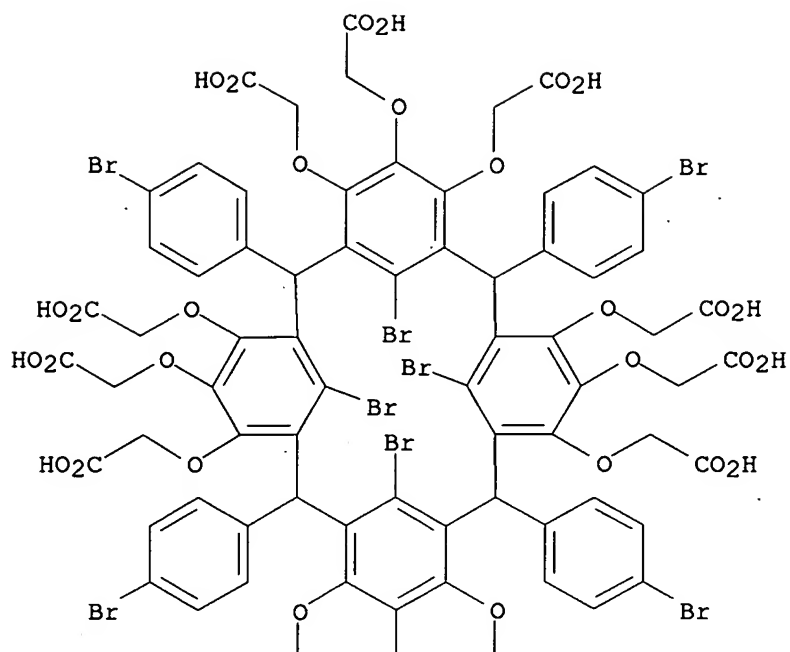
RN	171799-82-9	CAPLUS
CN	Acetic acid, 2,2',2'',2''',2'''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''',2'''''''''-[25,26,27,28-tetrabromo-2,8,14,20-(10-chloro- 9-anthracenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos- 1(25'),3,5,7(28),8,11,13(27),15,17,19(26),21,23-dodecaene- 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis-, dodecaammonium salt (9CI) (CA INDEX NAME)	





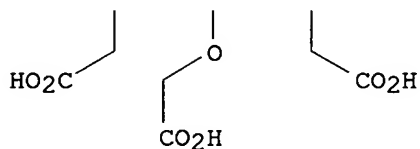
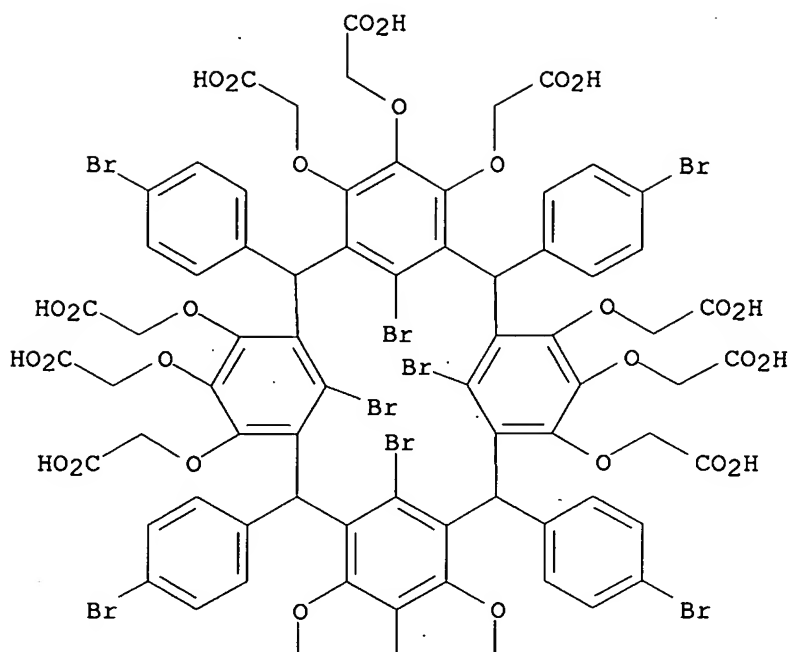
● 12 NH₃

RN	171799-89-6 CAPLUS
CN	Acetic acid, 2,2',2'',2''',2'''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''',2'''''''''-[25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4- bromophenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos- 1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene- 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl)dodecakis(oxy)]dodecakis-, dodecapotassium salt (9CI) (CA INDEX NAME)

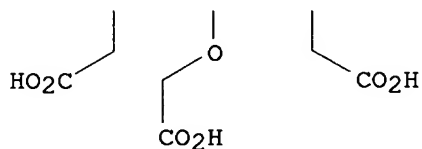
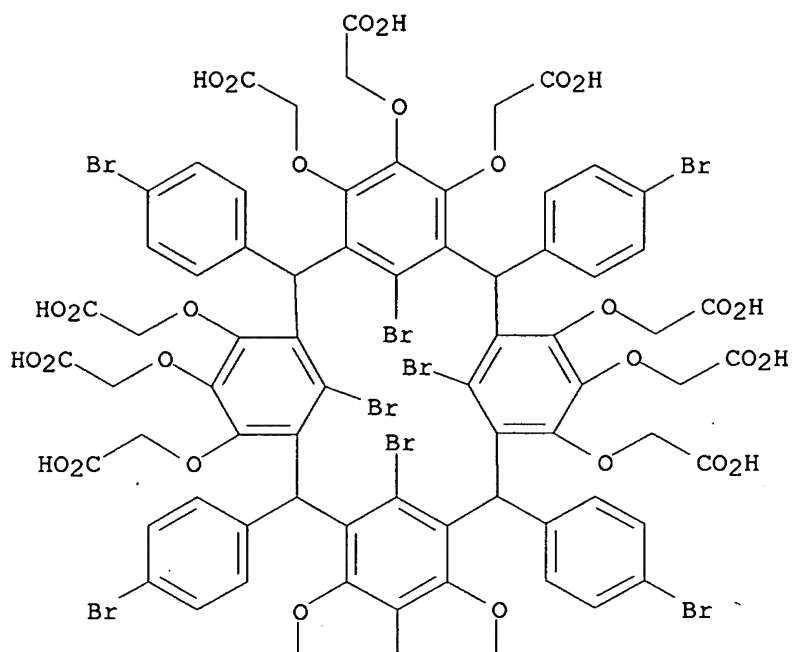


●12 K

RN	171799-90-9 CAPLUS
CN	Acetic acid, 2,2',2'',2''',2'''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''',2'''''''''-[25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4- bromophenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos- 1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene- 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis- (9CI) (CA INDEX NAME)

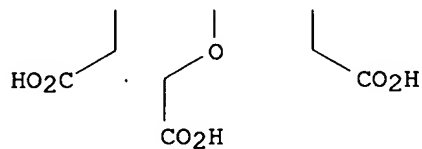
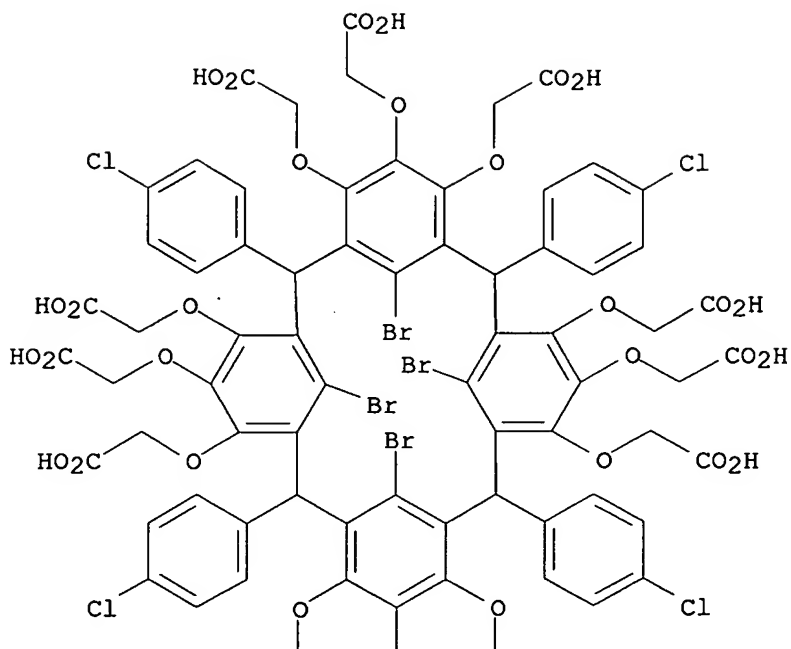


RN	171799-91-0	CAPLUS
CN	Acetic acid, 2,2',2'',2''',2'''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''''',2'''''''''''-[25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4- bromophenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos- 1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene- 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis-, dodecaammonium salt (9CI) (CA INDEX NAME)	



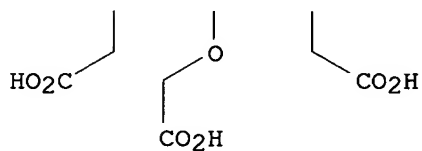
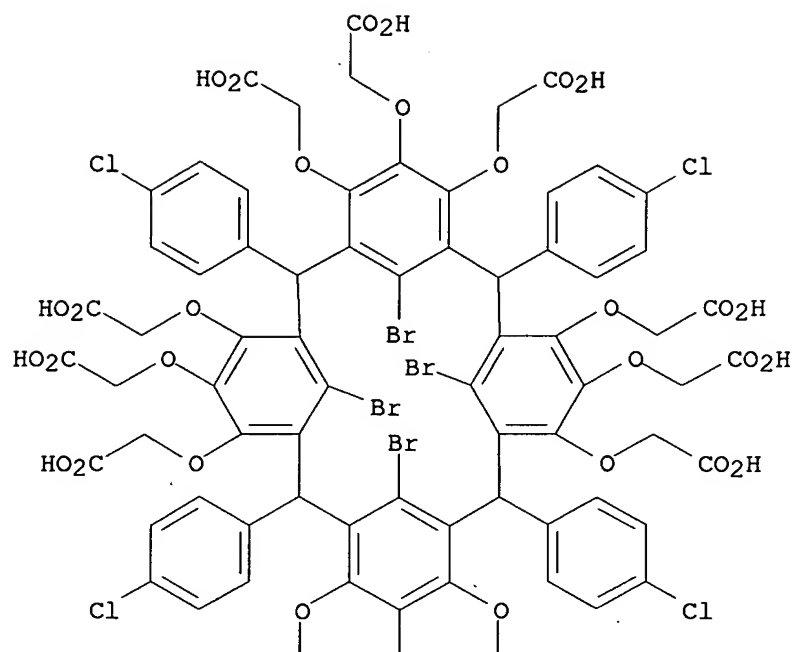
● 12 NH₃

RN	171799-95-4	CAPLUS
CN	Acetic acid, 2,2',2'',2''',2'''',2''''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''',2'''''''''-[25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4- chlorophenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos- 1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene- 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis-, dodecapotassium salt (9CI) (CA INDEX NAME)	

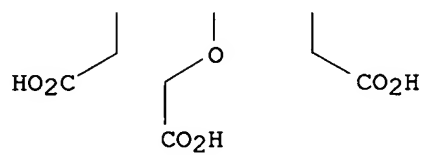
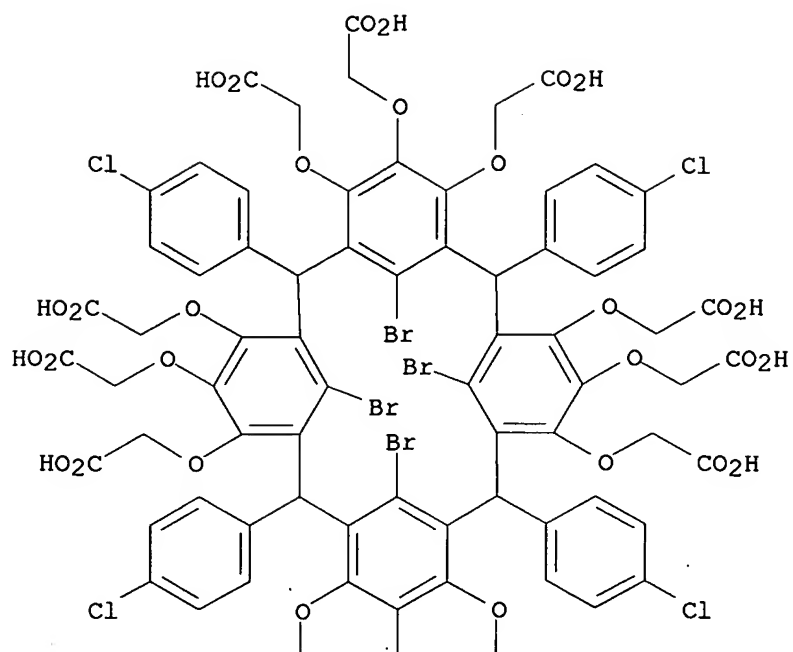


● 12 K

RN	171799-96-5	CAPLUS
CN	Acetic acid, 2,2',2'',2''',2''''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''''',2'''''''''''-[25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4- chlorophenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacos- 1(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene- 4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis- (9CI) (CA INDEX NAME)	



RN	171799-97-6	CAPLUS
CN	Acetic acid, 2,2',2'',2''',2'''',2''''',2''''',2''''',2''''',2''''', ''',2''''''''',2'''''''''-[[25,26,27,28-tetrabromo-2,8,14,20-tetrakis(4-chlorophenyl)pentacyclo[19.3.1.13,7.19,13.115,19]octacosal(25),3,5,7(28),9,11,13(27),15,17,19(26),21,23-dodecaene-4,5,6,10,11,12,16,17,18,22,23,24-dodecayl]dodecakis(oxy)]dodecakis-, didecaammonium salt (9CI) (CA INDEX NAME)	



●12 NH₃